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## WHAT IS CLAIMED IS:

- 1. A method for inducing T-cell tolerance or non-responsiveness of donor T-cells to desired alloantigen or xenoantigen bearing cells *in vitro* comprising the following:
  - (i) providing a culture containing donor tissue containing donor T-cells;
- (ii) producing a mixed lymphocyte reaction culture by adding to said donor T-cell culture alloantigen or xenoantigen-bearing cells;
- (iii) adding to the resultant mixed lymphocyte culture a gp39 antagonist; and
- 10 (iv) maintaining these cells in culture for a sufficient time to render the donor T-cells substantially non-responsiveness to said alloantigen or xenoantigen bearing cells.
  - 2. The method of Claim 1, wherein the tissue containing donor T-cells is donor bone marrow or peripheral blood cells.
    - 3. The method of Claim 1, wherein the gp39 antagonist is selected from the group consisting of an anti-gp39 antibody, soluble CD40 and soluble CD40 fusion protein.

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- 4. The method of Claim 3, wherein the gp39 antagonist is an anti-gp39 antibody.
- 5. The method of Claim 4, wherein said anti-gp39 antibody is an antibuman gp39 monoclonal antibody.
  - 6. The method of Claim 1, wherein the donor T-cells are cultured with said gp39 antagonist for a time ranging from about 1 to 30 days.
- The method of Claim 6, wherein said time ranges from 5 to 15 days.
  - 8. The method of Claim 1, wherein the alloantigen or xenoantigen bearing cells comprise cells or tissue obtained from a potential transplant recipient that has been treated to deplete recipient T-cells.
  - 9. The method of Claim 8, wherein T-cell depletion is effected by irradiation.

- 10. The method of Claim 1, wherein the donor T-cells are transplanted into a recipient in need of such transplantation.
- 11. The method of Claim 10, wherein the recipient is in need of immune reconstitution as a result of disease or disease treatment.
  - 12. The method of Claim 11, wherein said disease is cancer or autoimmune disease.